

Middlebrook 7H10 Agar



Medium used for the isolation and cultivation of Mycobacteria.

• CONTENTS (Liter)

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|------------------------------|-----------|
| Disodium Phosphate | 1.5 g |
| Monopotassium Phosphate | 1.5 g |
| Ammonium Sulfate | 0.5 g |
| L-Glutamic Acid | 0.5 g |
| Sodium Citrate | 0.4 g |
| Ferric Ammonium Citrate | 0.04 g |
| Magnesium Sulfate | 0.025 g |
| Copper Sulfate | 0.001 g |
| Pyridoxine Hydrochloride | 0.001 g |
| Zinc Sulfate | 0.001 g |
| Biotin | 0.0005 g |
| Calcium Chloride | 0.0005 g |
| Malachite Green | 0.00025 g |
| Agar | 15.0 g |
| Final pH = 6.6 ± 0.2 at 25°C | |

• PROCEDURE

Suspend 19.47 G of powder in 900 mL of distilled or deionized water. Add 5 mL of Glycerol supplement (MB-G1821). Mix well. Heat to boiling until completely dissolved. Sterilize by autoclave at 121°C for 15 minutes. Cool to 45 - 50°C in water bath. Aseptically add 2 vials of Middlebrook OADC Enrichment supplement (MB-M3021). Mix well. Pour into petri dishes.

Middlebrook OADC Enrichment supplement

1 vial contents (each vial is sufficient for 500 mL of medium)

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|----------------------------|----------|
| Oleic Acid | 0.025 mL |
| Albumin Fraction V, Bovine | 2.5 g |
| Glucose | 1.0 g |
| Catalase | 0.002 g |
| Sodium Chloride | 0.425 g |

• INTERPRETATION

Middlebrook 7H10 Agar is a medium used for the isolation and cultivation of Mycobacteria. Disodium phosphate, monopotassium phosphate, ammonium sulfate, L-glutamic acid, ferric ammonium citrate, magnesium sulfate, copper sulfate, pyridoxine hydrochloride, zinc sulfate, biotin and calcium chloride are inorganic salts essential for the growth of Mycobacteria. Sodium citrate is converted to citric acid which holds inorganic cations in solution. Malachite green serves as the selective agent to inhibit bacteria except Mycobacteria. Agar is the solidifying agent. Glycerol is the carbon and energy source. Oleic acid is necessary in the metabolism of Mycobacteria. Catalase destroys toxic peroxides that may be present in the medium. Albumin protects Mycobacteria against toxic agents. Glucose is the carbohydrate. Sodium chloride maintains the osmotic balance.

• TECHNIC

Inoculate the specimen using a sterile loop to the medium. Incubate at 35 ± 2°C for 2 - 5 days up to 3 weeks under microaerobic condition. Refer appropriate references for recommended test procedure.

• QUALITY CONTROL FOR USE

Dehydrated medium

Appearance: free-flowing, homogeneous

Color: light beige with greenish tint

Prepared medium

Appearance: clear to slightly opalescent

Color: greenish light amber

Incubation conditions: $35 \pm 2^{\circ}\text{C}$ / 2 - 5 days up to 3 weeks under microaerobic condition

| Microorganism | ATCC | Growth |
|--------------------------------|-------|-----------|
| <i>Mycobacterium smegmatis</i> | 607 | good |
| <i>Staphylococcus aureus</i> | 25923 | inhibited |

• STORE

The powder is very hygroscopic. Store the powder at room temperature, in a dry environment, in its original container tightly closed and use it before the expiry date on the label. Store prepared medium at $2 - 8^{\circ}\text{C}$.

• REFERENCES

1. Musser, J.M. 1995. Clin. Microbiol. Rev. 8: 496-514.
2. Kleitmann, W. 1995. Clin. Microbiol. News. 17: 65-69.
3. Middlebrook, G., M.L. Cohn, W.B. Dye, W.B. Russell, Jr., D. and Levy. 1960. Acta. Tubercul. Scand. 38: 66.

• PACKAGE

| | |
|---|-------|
| Cat. No : MB-M1919 Middlebrook 7H10 Agar | 500 G |
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